

R2
Sub C2
5. (Amended) A fast-dissolving pharmaceutical composition, which comprises micronized AS-3201 in a ratio of about 0.5% by weight - 5% by weight, a diluent in a ratio of about 51% by weight - about 93.8% by weight, a disintegrator in a ratio of about 5% by weight - about 35% by weight, a binder in a ratio of about 0.5% by weight - about 5% by weight, and a lubricant in a ratio of about 0.2% by weight - about 4% by weight, relative to the total weight of the pharmaceutical composition,

wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 50% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

Sub C3
R3
13. (Amended) A fast-dissolving pharmaceutical composition, which comprises micronized AS-3201 in a ratio of more than 5% by weight and less than about 25% by weight, a diluent in a ratio of about 16% by weight - about 84.3% by weight, a disintegrator in a ratio of about 10% by weight - about 50 % by weight, a binder in a ratio of about 0.5% by weight - about 5% by weight, and a lubricant in a ratio of about 0.2 % by weight - about 4% by weight, relative to the total weight of the pharmaceutical composition,

wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 50% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

Please add the following new claims:

Sub C4
R4
63. (New) The fast-type dissolving pharmaceutical composition according to claim 1, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

64. (New) The fast-type dissolving pharmaceutical composition according to claim 2, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

65. (New) The fast-type dissolving pharmaceutical composition according to claim 3, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

66. (New) The fast-type dissolving pharmaceutical composition according to claim 4, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

67. (New) The fast-type dissolving pharmaceutical composition according to claim 5, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

68. (New) The fast-type dissolving pharmaceutical composition according to claim 6, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

69. (New) The fast-type dissolving pharmaceutical composition according to claim 7, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-

3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

70. (New) The fast-type dissolving pharmaceutical composition according to claim 8, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

71. (New) The fast-type dissolving pharmaceutical composition according to claim 9, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

72. (New) The fast-type dissolving pharmaceutical composition according to claim 10, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

73. (New) The fast-type dissolving pharmaceutical composition according to claim 11, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

74. (New) The fast-type dissolving pharmaceutical composition according to claim 12, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

75. (New) The fast-type dissolving pharmaceutical composition according to claim 13, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

76. (New) The fast-type dissolving pharmaceutical composition according to claim 14, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

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77. (New) The fast-type dissolving pharmaceutical composition according to claim 15, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

78. (New) The fast-type dissolving pharmaceutical composition according to claim 16, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

79. (New) The fast-type dissolving pharmaceutical composition according to claim 17, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

80. (New) The fast-type dissolving pharmaceutical composition according to claim 18, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-

3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

Sub C4 Contd
81. (New) The fast-type dissolving pharmaceutical composition according to claim 19, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.

82. (New) The fast-type dissolving pharmaceutical composition according to claim 20, wherein when said composition is compressed into a tablet and a dissolution percentage of AS-3201 from the tablet is measured according to the Paddle method, 80% or more of the AS-3201 in the tablet is dissolved within 15 minutes from the start of the method.